In the Claims

1. (Currently Amended) A therapeutic or prophylactic agent for method of treating urinary frequency frequency, urinary urgency or urinary incontinence, comprising as an effective ingredient administering a therapeutically effective amount of a morphinan derivative having a nitrogen-containing heterocyclic group of the Formula (I):

$$R^{1}$$
 R^{2}
 R^{10}
 R^{11}
 R^{3}
 R^{3}
(I)

[[([]]wherein R^1 is hydrogen, C_1 - C_5 alkyl, C_4 - C_7 cycloalkylalkyl, C_6 - C_8 cycloalkenylalkyl, C_6 - C_{12} aryl, C_7 - C_{13} aralkyl, C_3 - C_7 alkenyl, furanylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5), thienylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5) or pyridylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5), R^2 and R^3 independently are hydrogen, hydroxy, C_1 - C_5 alkoxy, C_3 - C_7 alkenyloxy, C_7 - C_{13} aralkyloxy or C_1 - C_5 alkanoyloxy; Y and Z independently represent valence bond or -C(=O)-; -X- represents a C_2 - C_7 carbon chain (one or more of the carbon atoms therein m a y be substituted replaced by nitrogen, oxygen or sulfur atom(s), and the carbon chain may contain an unsaturated bond) constituting a part of the ring structure; k is an integer of 0 to 8; R^4 is(are) (a) substituent(s) in the number of k on the nitrogen-containing ring, which independently represent(s) fluorine, chlorine, bromine, iodine, nitro, hydroxy, C_1 - C_5 alkyl, benzylidene, ethylidene, cyclohexylmethylidene, butylidene, phenethylidene, C_7 - C_{13} cycloalkylalkyl, C_6 - C_{12} aryl, C_7 - C_{13} aralkyl, C_7 - C_{13} aralkyloxy, C_1 - C_5 alkoxy, trifluoromethyl, trifluoromethoxy, cyano, isothiocyanato, SR^6 , SOR^6 , SO_2R^6 , $(CH_2)_0OR^6$,

 $(CH_2)_pCOR^6$, $(CH_2)_pCO_2R^6$, $SO_2NR^7R^8$, $CONR^7R^8$, $(CH_2)_pNR^7R^8$ or $(CH_2)_pN(R^7)COR^8$, or among the R4s in the number of k, two R4s bound to the same carbon atom or to the same sulfur atom cooperatively represent one oxygen atom to form carbonyl or sulfoxide (with the proviso that in cases where Y and Z is a valence bond, the formed carbonyl is not bound directly to the nitrogen atom which is bound to the morphinan structure), or two R⁴s bound to the same carbon atom cooperatively represent one sulfur atom to form thiocarbonyl, or four R⁴s bound to the same sulfur atom cooperatively represent two oxygen atoms to form sulfone, or among the R⁴s in the number of k, two R⁴s bound to adjacent carbon atoms, respectively, cooperatively form benzene condensed fused ring, pyridine condensed fused ring, naphthalene condensed fused ring, cyclopropane fused ring, cyclobutane fused ring, cyclopentane fused ring, cyclopentene fused ring, cyclohexane fused ring, cyclohexene fused ring, cycloheptane fused ring or cycloheptene fused ring, each of said fused rings is non-substituted or substituted by 1 or more R⁵s, wherein R⁵(s) independently represent(s) fluorine, chlorine, bromine, iodine, nitro, hydroxy, C₁-C₅ alkyl, C₇-C₁₃ aralkyl (in cases where Y and Z are simultaneously -C(=0)- or valence bonds), C_1 - C_5 alkoxy, trifluoromethyl, trifluoro-methoxy, cyano, C₆-C₁₂ aryl, isothiocyanato, SR⁶, SOR⁶, SO₂R⁶, (CH₂)_pOR⁶, (CH₂)_pCOR⁶, (CH₂)_pCO₂R⁶, SO₂NR⁷R⁸, CONR⁷R⁸, (CH₂)_pNR⁷R⁸ or (CH₂)pN(R⁷)COR⁸; R⁹ is hydrogen, C₁-C₅ alkyl, C₁-C₅ alkenyl, C₇-C₁₃ aralkyl, C₁-C₃ hydroxyalkyl, (CH₂)pOR⁶ or (CH₂)pCO₂R⁶; R¹⁰ and R¹¹ are bound to form -O-, -S- or -CH₂-, or R¹⁰ is hydrogen and R¹¹ is hydrogen, hydroxy, C₁-C₅ alkoxy or C₁-C₅ alkanoyloxy; p is an integer of 0 to 5; R⁶ is hydrogen, C₁-C₅ alkyl, C₃-C₇ alkenyl, C₆-C₁₂ aryl or C_7 - C_{13} aralkyl; and R^7 and R^8 independently are hydrogen, C_1 - C_5 alkyl or C_7 - C_{13} aralkyl[[]]; or a pharmaceutically acceptable acid addition salt thereof to a patient.

2. (Currently Amended) The therapeutic or prophylactic agent method according to claim 1, wherein in said Formula (I), only one of Y and Z is -C(=O)- and the other is valence bond.

- 3. (Currently Amended) The therapeutic or prophylactic agent method according to claim 1, wherein in said Formula (I), both Y and Z are -C(=O)-.
- 4. (Currently Amended) The therapeutic or prophylactic agent method according to claim 3, wherein in said Formula (I), R¹ is hydrogen, C₄-C₇ cycloalkylalkyl, C₆-C₈ cycloalkenylalkyl, C₆-C₁₂ aryl or C₃-C₇ alkenyl; k is an integer of 2 to 8; and two R⁴s bound to adjacent carbon atoms, respectively, cooperatively form benzene fused ring, pyridine fused ring, naphthalene fused ring, cyclopropane fused ring, cyclobutane fused ring, cyclopentane fused ring, cyclopentene fused ring, cyclohexane fused ring, cyclohexene fused ring, cycloheptane fused ring or cycloheptene fused ring, each of said fused rings is non-substituted or substituted by 1 or more R⁵s.
- 5. (Currently Amended) The therapeutic or prophylactic agent method according to claim 3, wherein in said Formula (I), R¹ is hydrogen, cyclopropylmethyl, cyclobutylmethyl, cyclobexylmethyl, allyl or prenyl; R² is hydrogen, hydroxy, methoxy, ethoxy, allyloxy, benzyloxy, acetoxy or propionoxy; R³ is hydrogen, hydroxy, methoxy, ethoxy, benzyloxy, acetoxy or propionoxy; k is an integer of [[0]]2 to 6, two R⁴s cooperatively form benzene fused ring which is non-substituted or substituted by 1 to 4 R⁵s; R⁵(s) independently is(are) fluorine, chlorine, bromine, iodine, nitro, methyl, ethyl, propyl, benzyl, hydroxy, methoxy, ethoxy, trifluoromethyl, trifluoromethoxy, cyano, phenyl, isothiocyanato, SR⁶, SOR⁶, SO₂R⁶, (CH₂)_pOR⁶, (CH₂)_pCOR⁶, (CH₂)_pCO₂R⁶, SO₂NR⁷R⁸, CONR⁷R⁸, (CH₂)_pNR⁷R⁸ or (CH₂)_pN(R⁷)COR⁸; p is an integer of 0 to 5; R⁶ is hydrogen, methyl, ethyl, propyl or phenyl; R⁷ and R⁸ independently are hydrogen, methyl, ethyl, propyl or benzyl; R⁹ is hydrogen or methyl; R¹⁰ and Rⁿ are bound to form -O-, or R¹⁰ is hydrogen and R¹¹ is hydrogen, hydroxy or methoxy.
 - 6. (Currently Amended) The therapeutic or prophylactic agent method according to

claim 1, wherein in said Formula (I), both Y and Z are valence bonds.

- 7. (Currently Amended) The therapeutic or prophylactic agent method according to claim 6, wherein in said Formula (I), R¹ is hydrogen, C₁-C₅ alkyl, C₇-C₁₃ aralkyl, furanylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5), thienylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5) or pyridylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5); k is an integer of 2 to 8; two R⁴s bound to adjacent carbon atoms, respectively, cooperatively form benzene fused ring, pyridine fused ring, naphthalene fused ring, cyclopropane fused ring, cyclobutane fused ring, cyclopentane fused ring, cyclopentene fused ring, cyclohexane fused ring, cyclohexene fused ring, cycloheptane fused ring or cycloheptene fused ring, each of said fused rings is non-substituted or substituted by 1 or more R⁵s.
- 8. (Currently Amended) The therapeutic or prophylactic agent method according to claim 6, wherein in said Formula (I), R¹ is hydrogen, methyl, ethyl, propyl, benzyl, phenethyl, phenylpropyl, 2-furanylmethyl, 2-furanylethyl, 2-furanylpropyl, 3-furanylmethyl, 3-furanylethyl, 3furanylprofyl, 2-thiophenylmethyl 2-thionylmethyl, 2-thiophenylethyl 2-thionylethyl, 2thiophenylpropyl 2-thiophenylmethyl 3-thiophenylmethyl 3-thiophenylethyl 3thienylethyl, 3-thiophenylpropyl 3-thienylpropyl, 2-pyridynylmethyl, 2-pyridynylethyl, 2-pyridynylpropyl, 3-pyridynylmethyl, 3-pyridynylpropyl, 4-pyridynylmethyl, 4pyridynylethyl, or 4-pyridynylpropyl; R² is hydrogen, hydroxy, methoxy, ethoxy, allyloxy, benzyloxy, acetoxy or propionoxy; R³ is hydrogen, hydroxy, methoxy, ethoxy, benzyloxy, acetoxy or propionoxy; k is an integer of [[0]]2 to 6; two R⁴s cooperatively form benzene fused ring which is non-substituted or substituted by 1 to 4 R⁵s and other R⁴(s) independently is(are) methyl, ethyl, propyl or benzyl, or two R⁴s bound to the same carbon atom represent one oxygen atom to form carbonyl, R⁵(s) independently is(are) fluorine, chlorine, bromine, iodine, nitro, methyl, ethyl, propyl,

benzyl, hydroxy, methoxy, ethoxy, trifluoromethyl, trifluoromethoxy, cyano, phenyl, isothiocyanato, SR^6 , SOR^6 , SO_2R^6 , $(CH_2)_pOR^6$, $(CH_2)_pCOR^6$, $(CH_2)_pCO_2R^6$, $SO_2NR^7R^8$, $CONR^7R^8$, $(CH_2)_pNR^7R^8$ or $(CH_2)pN(R^7)COR^8$; p is an integer of 0 to 5; R^6 is hydrogen, methyl, ethyl, propyl or phenyl; R^7 and R^8 independently are hydrogen, methyl, ethyl, propyl or benzyl; R^9 is hydrogen or methyl; R^{10} and R^{11} are bound to form -O-, or R^{10} is hydrogen and R^{11} is hydrogen, hydroxy or methoxy.

- 9. (Cancelled)
- 10. (Cancelled)
- 11. (Currently Amended) A morphinan derivative of the Formula (II) having a nitrogencontaining heterocyclic group:

[[(]]wherein R^1 , R^2 , R^3 , R^9 , R^{10} and R^{11} represent are the same meanings as described above in claim 1, R^4 , X, Y, Z and R^4 , R^4

or a pharmaceutically acceptable acid addition salt thereof.

- 12. (Original) The morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 11, wherein in said Formula (II), only one of Y' and Z' is C(=O)- and the other is valence bond.
- 13. (Original) The morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 11, wherein in said Formula (II), both Y' and Z' are C(=O)-.
- 14. (Currently Amended) The morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 13, wherein in said Formula (II), R¹ is hydrogen, [[C₄-C₇]] C₄-C₇ cycloalkylalkyl, C₆-C₈ cycloalkenylalkyl, C₆-C₁₂ aryl or [[C₃-C₇]] C₃-C₇ alkenyl; k' is an integer of 2 to 8 and two R⁴s bound to adjacent carbon atoms, respectively, cooperatively form benzene fused ring substituted by 1 or more R⁵s, or cooperatively form a pyridine fused ring, naphthalene fused ring, cyclopropane fused ring, cyclobutane fused ring, cyclopentane fused ring or cyclopentene fused ring, cyclohexane fused ring, cyclohexene fused ring, cycloheptane fused ring or cycloheptene fused ring, each of said fused rings is non-substituted or substituted by 1 or more R⁵s.
- 15. (Currently Amended) The morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 13, wherein in said Formula (II), R¹ is hydrogen, cyclopropylmethyl, cyclobutylmethyl, cyclopentylmethyl, cyclohexylmethyl, allyl or prenyl; R² is hydrogen, hydroxy, methoxy, ethoxy, allyloxy, benzyloxy, acetoxy or propionoxy; R³ is hydrogen, hydroxy, methoxy, ethoxy, benzyloxy, acetoxy or propionoxy; k' is an integer of [[0]]2 to 6, two R⁴'s cooperatively form benzene fused ring which is non-substituted or substituted by 1 to 4 R⁵s; R⁵(s) independently is(are) fluorine, chlorine, bromine, iodine, nitro, methyl, ethyl, propyl, benzyl, hydroxy, methoxy, ethoxy, trifluoromethyl, trifluoromethoxy, cyano, phenyl, isothiocvanato, SR⁶,

SOR⁶, SO₂R⁶, (CH₂)_POR⁶, (CH₂)_PCOR⁶, (CH₂)_PCO₂R⁶, SO₂NR⁷R⁸, CONR⁷R⁸, (CH₂)_PNR⁷R⁸ or $(CH_2)_PN(R^7)COR^8$; p is an integer of 0 to 5; R⁶ is hydrogen, methyl, ethyl, propyl or phenyl; R⁷ and R⁸ independently are hydrogen, methyl, ethyl, propyl or benzyl; R⁹ is hydrogen or methyl; R¹⁰ and R¹¹ are bound to form -O-, or R¹⁰ is hydrogen and R¹¹ is hydrogen, hydroxy or methoxy.

- 16. (Original) The morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 11, wherein in said Formula (II), both Y' and Z' are valence bonds.
- 17. (Currently Amended) The morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 16, wherein in said Formula (II), R¹ is hydrogen, C₁-C₅ alkyl, C₇-C₁₃ aralkyl, furanylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5), thienylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5) or pyridylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5); k' is an integer of 6 to 8; two R⁴'s bound to adjacent carbon atoms, respectively, cooperatively form benzene fused ring, pyridine fused ring, naphthalene fused ring, cycloperopane fused ring, cyclobutane fused ring, cyclopentane fused ring, cyclopentene fused ring, cyclohexane fused ring, cyclohexene fused ring, cycloheptane fused ring or cycloheptene fused ring, each of said fused rings is non-substituted or substituted by 1 or more R⁵s.
- 18. (Currently Amended) The morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 16, wherein in said Formula (II), R¹ is hydrogen, methyl, ethyl, propyl, benzyl, phenethyl, phenylpropyl, 2-furanylmethyl, 2-furanylethyl, 2-furanylpropyl, 3-furanylmethyl, 3-furanylpropyl, 2-thiophenylmethyl 2-thiophenylethyl, 2-thiophenylpropyl, 2-thiophenylpropyl, 3-thiophenylpropyl, 3-thiophenylpropyl, 3-thiophenylpropyl, 3-thiophenylpropyl, 2-thiophenylpropyl, 3-thiophenylpropyl, 3-thio

pyridynylmethyl, 2-pyridynylethyl, 2-pyridynylpropyl, 3-pyridynylmethyl, 3-pyridynylpropyl, 4-pyridynylmethyl, 4-pyridynylpropyl, R² is hydrogen, hydroxy, methoxy, ethoxy, allyloxy, benzyloxy, acetoxy or propionoxy; R³ is hydrogen, hydroxy, methoxy, benzyloxy, acetoxy or propionoxy; k' = 6 [[is an integer of 0 to 6]]; two R⁴'s cooperatively form benzene fused ring which is non-substituted or substituted by 1 to 4 R⁵s and other R⁴(s) independently is(are) methyl, ethyl, propyl or benzyl, or two R⁴'s bound to the same earbon atom represent one oxygen atom to form carbonyl, R⁵(s) independently is(are) fluorine, chlorine, bromine, iodine, nitro, methyl, ethyl, propyl, benzyl, hydroxy, methoxy, ethoxy, trifluoromethyl, trifluoromethoxy, cyano, phenyl, isothiocyanato, SR⁶, SOR⁶, SO₂R⁶, (CH₂)pOR⁶, (CH₂)pCO₂R⁶, SO₂NR⁷R⁸, CONR⁷R⁸, (CH₂)pNR⁷R⁸ or (CH₂)pN(R⁷)COR⁸; p is an integer of 0 to 5; R⁶ is hydrogen, methyl, ethyl, propyl or phenyl; R⁷ and R⁸ independently are hydrogen, methyl, ethyl, propyl or benzyl; R⁹ is hydrogen or methyl; R¹⁰ and R¹¹ are bound to form -O-, or R¹⁰ is hydrogen and R¹¹ is hydrogen, hydroxy or methoxy.

- 19. (Cancelled)
- 20. (Previously Presented) A pharmaceutical composition comprising the morphinan derivative or the pharmaceutically acceptable acid addition salt thereof according to claim 11.
 - 21. (Cancelled)
 - 22. (Cancelled)
 - 23. (Cancelled)